

# Draft Genome Sequence of *Xanthomonas sacchari* Strain LMG 476

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**We report the high-quality draft genome sequence of *Xanthomonas sacchari* strain LMG 476, isolated from sugarcane. The genome comparison of this strain with a previously sequenced *X. sacchari* strain isolated from a distinct environmental source should provide further insights into the adaptation of this species to different habitats and its evolution.**

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To date, the *Xanthomonas* genus accounts for 37 plant-associated species and subspecies according to the current prokaryotic nomenclature listing at the Leibniz Institute DSMZ (<http://www.dsmz.de/bacterial-diversity/prokaryotic-nomenclature-up-to-date.html>). However, the recently proposed new species *X. maliensis* isolated from rice could be added to this list (1). The xanthomonads' taxonomy evolves regularly according to the methodologies used for description, resulting in changes in the distribution and delimitation of species, as well as in the reallocation of pathovars within species. In 1995, on the basis of DNA-DNA hybridization analyses, 20 *Xanthomonas* species (genomospecies) were proposed, elevating many *X. campestris* pathovars to the rank of species or placing them in already described species (2). This work resulted also in the repositioning of strains LMG 471 and LMG 476, isolated from sugarcane in Guadeloupe in 1980. These two strains, initially considered to belong to the sugarcane pathogenic species *X. albilineans*, were renamed as a new *Xanthomonas* species called *X. sacchari* (2). Thereafter, phylogenetic trees based on 16S ribosomal DNA or 16S-23S ribosomal DNA intergenic sequences, as well as multilocus sequence analyses, supported the distinction of both LMG 471 and LMG 476 strains from the *X. albilineans* species, nevertheless confirming that the *X. sacchari* species is genetically close to *X. albilineans* (3–5). *X. albilineans* is a sugarcane pathogen causing leaf scald disease, whereas the pathogenicity of *X. sacchari* remains unknown. An additional *X. sacchari* strain, NCPPB4393, isolated from an insect collected on a banana plant in Tanzania in 2007, has recently been sequenced (6, 7). The lack of information regarding both the adaptation of *X. sacchari* to two different habitats (insect and plant) and the specific genomic traits of this species raised the interest to compare strain NCPPB4393 with an *X. sacchari* LMG strain. We report herein the draft genome sequence of *X. sacchari* strain LMG 476, which is considered the type strain of this species (2).

Strain LMG 476 was sequenced using a Solexa HiSeq (Illumina) sequencer (Genoscope, France). The shotgun sequencing yielded 72,261,076 76-bp paired-end reads with an insert size of 200 bp. A combination of Velvet (8), SOAPdenovo, and SOAP-

GapCloser (9) yielded 74 contigs larger than 500 bp and a largest scaffold of 4,597,345 bp (using *X. albilineans* strain GPE PC73 as a reference for guiding the final scaffolding step) for a total assembly size of 4,898,860 bp (excluding scaffolding gaps), corresponding to 1,121× coverage. The average G+C content is 69%.

Calculation of the average nucleotide identity (ANI) was performed between the genome sequences of strain LMG 476, strain NCPPB4393, and strain GPE PC73 of *X. albilineans* using the JSpecies calculator (<http://www.imedeia.uib.es/jspecies>) (10) based on the BLASTn method (ANIb) and the MUMer algorithm (ANIm). The ANI values between strains LMG 476 and NCPPB4393 were above 99%, whereas those between strains LMG 476 or NCPPB4393 and GPE PC73 were below 87%, thus confirming that strain NCPPB4393 belongs to the *X. sacchari* species. The genome sequence of strain LMG 476 can now be used to elucidate the genomics and evolution of this species of *Xanthomonas*, whose habitats remain unclear, especially because strain NCPPB4393 was isolated from an insect.

**Nucleotide sequence accession numbers.** This whole-genome shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession number [JXQE000000000](https://www.ncbi.nlm.nih.gov/nuccore/JXQE000000000). The version described in this paper is version [JXQE010000000](https://www.ncbi.nlm.nih.gov/nuccore/JXQE010000000).

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## REFERENCES

1. Triplett LR, Verdier V, Campillo T, Van Malderghem C, Cleenwerck I, Maes M, Deblais L, Corral R, Koita O, Cottyn B, Leach JE. 15 January 2015. Characterization of a novel clade of *Xanthomonas* isolated from rice leaves in Mali and proposal of *Xanthomonas maliensis* sp. nov. *Antonie Van Leeuwenhoek*. <http://dx.doi.org/10.1007/s10482-015-0379-5>.
2. Vauterin L, Hoste B, Kersters K, Swings J. 1995. Reclassification of *Xanthomonas*. *Int J Syst Bacteriol* 45:472–489. <http://dx.doi.org/10.1099/00207713-45-3-472>.
3. Hauben L, Vauterin L, Swings J, Moore ER. 1997. Comparison of 16S ribosomal DNA sequences of all *Xanthomonas* species. *Int J Syst Bacteriol* 47:328–335. <http://dx.doi.org/10.1099/00207713-47-2-328>.
4. Gonçalves ER, Rosato YB. 2002. Phylogenetic analysis of *Xanthomonas*

- species based upon 16S-23S rDNA intergenic spacer sequences. *Int J Syst Evol Microbiol* 52:355–361. <http://dx.doi.org/10.1099/ijs.0.01886-0>.
5. Young JM, Park DC, Shearman HM, Fargier E. 2008. A multilocus sequence analysis of the genus *Xanthomonas*. *Syst Appl Microbiol* 31: 366–377. <http://dx.doi.org/10.1016/j.syapm.2008.06.004>.
  6. Studholme DJ, Wasukira A, Paszkiewicz K, Aritua V, Thwaites R, Smith J, Grant M. 2011. Draft genome sequences of *Xanthomonas sacchari* and two banana-associated xanthomonads reveal insights into the *Xanthomonas* group 1 clade. *Genes* 2:1050–1065. <http://dx.doi.org/10.3390/genes2041050>.
  7. Studholme DJ, Wasukira A, Paszkiewicz K, Aritua V, Thwaites R, Smith J, Grant M. 2012. Correction: Studholme *et al.*, draft genome sequences of *Xanthomonas sacchari* and two banana-associated xanthomonads reveal insights into the *Xanthomonas* group 1 clade. *Genes* 2011, 2, 1050–1065. *Genes* 3:88–89. <http://dx.doi.org/10.3390/genes3010088>.
  8. Zerbino DR, Birney E. 2008. Velvet: algorithms for *de novo* short read assembly using de Bruijn graphs. *Genome Res* 18:821–829. <http://dx.doi.org/10.1101/gr.074492.107>.
  9. Luo R, Liu B, Xie Y, Li Z, Huang W, Yuan J, He G, Chen Y, Pan Q, Liu Y, Tang J, Wu G, Zhang H, Shi Y, Liu Y, Yu C, Wang B, Lu Y, Han C, Cheung DW, Yiu SM, Peng S, Xiaoqian Z, Liu G, Liao X, Li Y, Yang H, Wang J, Lam TW, Wang J. 2012. SOAPdenovo2: an empirically improved memory-efficient short-read *de novo* assembler. *GigaScience* 1:18. <http://dx.doi.org/10.1186/2047-217X-1-18>.
  10. Richter M, Rosselló-Móra R. 2009. Shifting the genomic gold standard for the prokaryotic species definition. *Proc Natl Acad Sci U S A* 106: 19126–19131. <http://dx.doi.org/10.1073/pnas.0906412106>.